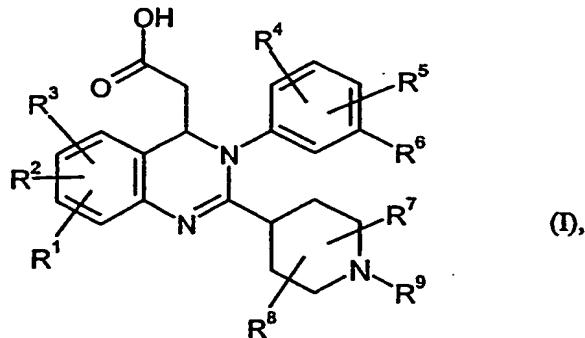


Patent Claims

1. Compound of the formula



5 in which

R¹, R² and R³ independently of one another represent hydrogen, alkyl, alkoxy, carboxyl, alkylcarbonyl, alkoxy carbonyl, aminocarbonyl, trifluoromethyl, halogen, cyano, hydroxyl or nitro,

10

R⁴ and R⁵ independently of one another represent hydrogen, alkyl, alkoxy, alkylthio, cyano, halogen, nitro, trifluoromethyl or trifluoromethoxy,

15

R⁶ represents alkyl, cyano, halogen, nitro or trifluoromethyl,

R⁷ and R⁸ independently of one another represent hydrogen, halogen, alkyl or alkoxy and

20

R⁹ represents aryl or 1,3-benzodioxol-5-yl, where aryl and 1,3-benzodioxol-5-yl may be substituted by 1 to 3 substituents, where the substituents independently of one another are selected from the group consisting of alkoxy, alkylthio, carboxyl, alkylcarbonyl, alkoxy carbonyl, aminocarbonyl, trifluoromethyl, halogen, carbamoyl, cyano, hydroxyl, amino,

25

alkylamino, nitro and optionally hydroxyl-substituted alkyl,

or one of its salts, its solvates or the solvates of its salts.

5 2. Compound according to Claim 1, characterized in that

R¹, R² and R³ independently of one another represent hydrogen, methyl, fluorine, chlorine, cyano, hydroxyl or aminocarbonyl,

10 R⁴ and R⁵ independently of one another represent hydrogen, fluorine, C₁-C₄-alkyl or C₁-C₄-alkoxy,

R⁶ represents chlorine, nitro, trifluoromethyl, methyl, isopropyl or tert-butyl,

15 R⁷ and R⁸ independently of one another represent hydrogen or C₁-C₃-alkyl and

20 R⁹ represents phenyl or 1,3-benzodioxol-5-yl, where phenyl may be substituted by 1 to 3 substituents, where the substituents independently of one another are selected from the group consisting of C₁-C₆-alkyl, C₁-C₆-alkoxy, carboxyl, C₁-C₆-alkylcarbonyl, C₁-C₆-alkoxycarbonyl, trifluoromethyl, fluorine, chlorine, bromine, cyano, hydroxyl, amino, C₁-C₆-alkylamino and nitro.

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3. Compound according to Claim 1 or 2, characterized in that

R¹ and R² are hydrogen,

30 R³ is fluorine,

R⁴ and R⁵ independently of one another are hydrogen, fluorine or

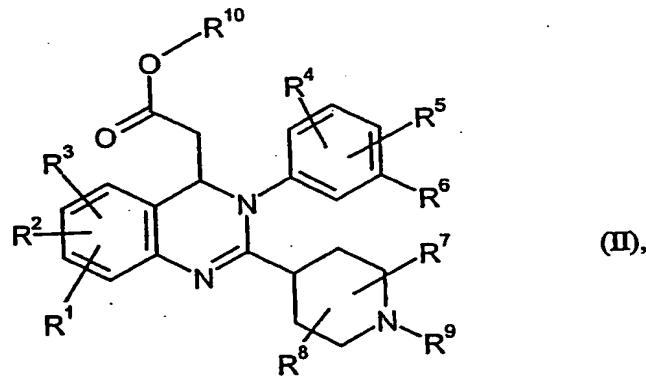
methoxy,

R⁶ is trifluoromethyl,

5 R⁷ and R⁸ are hydrogen and

10 R⁹ is phenyl, where phenyl may be substituted by 1 or 2 substituents, where the substituents independently of one another are selected from the group consisting of methyl, methoxy, ethoxy, fluorine and chlorine.

4. Process for preparing a compound of the formula (I) according to Claim 1, characterized in that a compound of the formula



15 in which

R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸ and R⁹ are as defined in Claim 1, and

R¹⁰ represents alkyl, preferably methyl or ethyl,

20 is reacted with a base.

5. Compound according to any of Claims 1 to 3 for the treatment and/or prophylaxis of diseases.

6. Use of a compound according to any of Claims 1 to 3 for preparing a medicament for the treatment and/or prophylaxis of diseases.
7. Use of a compound according to any of Claims 1 to 3 for preparing a medicament for the treatment and/or prophylaxis of viral infections.
8. Use according to Claim 7, characterized in that the viral infection is an infection with the human cytomegalovirus (HCMV) or another representative of the group of the Herpesviridae.
- 10 9. Medicament, comprising a compound as defined in any of Claims 1 to 3 in combination with a further active compound.
- 15 10. Medicament, comprising a compound according to any of Claims 1 to 3 in combination with an inert nontoxic, pharmaceutically acceptable auxiliary.
11. Medicament according to Claim 10 for the treatment and/or prophylaxis of viral infections.
- 20 12. Method for controlling viral infections in humans and animals by administering an antivirally effective amount of at least one compound according to any of Claims 1 to 3, a medicament according to any of Claims 9 to 11 or a medicament obtained according to any of Claims 6 to 8.